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 7/K/1 (Item 1 from file: 5) Links
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Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rights reserved. 0019555755 Biosis No.: 200700215496
0019555755
A panel of monoclonal antibodies recognizing the Staphylococcus epidermidis
fibrinogen-binding MSCRAMM SdrG
Author: Hall Andrea E; Patel Pratiksha R; Domanski Paul J; Prater Bradley D;
Gorovits Elena L; Syribeys Peter J; Vernachio John H; Patti Joseph M; Hutchins Jeff
Author Address: Inhibitex Inc, 9005 Westside Pkwy, Alpharetta, GA 30004 USA**USA
Author E-mail Address: jhutchins@inhibitex.com
Journal: Hybridoma 26 (1): p 28-34 FEB 2007 2007
ISSN: 1554-0014
Document Type: Article
Record Type: Abstract
Language: English
A panel of monoclonal antibodies recognizing the Staphylococcus epidermidis
fibrinogen-binding MSCRAMM SdrG
Abstract: ...stage contributing to the pathogenesis of this bacteria is the initial adherence to host tissue. SdrG is a cell-wall-anchored fibrinogen-binding adhesin of S. epidermidis that has been shown to be necessary for bacterial binding to fibrinogen-coated foreign bodies, such as catheters. Here we report the generation and characterization of a panel of monoclonal antibodies (MAbs) directed against this S. epidermidis virulence factor. Through the use of multiple in....that may prove to be beneficial in studies that address the precise biologic role of SdrG
prove to be beneficial in studies that address the precise biologic role of SdrG.
DESCRIPTORS:
 Chemicals & Biochemicals:
                                           fibrinogen; ... ...monoclonal antibody... ... SdrG
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    Fulltext available through: American Society for Microbiology custom link
USPTO Full Text Retrieval Options
Biosis Previews(R)
(c) 2007 The Thomson Corporation. All rights reserved. 18943764 Biosis No.: 200600289159
18943764
Human immunoglobulin G recognizing fibrinogen-binding surface proteins is protective against both Staphylococcus aureus and Staphylococcus epidermidis infections in vivo
Author: Vernachio John H (Reprint); Bayer Arnold S; Ames Brenda; Bryant Dawn; Prater
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Bradley D; Syribeys Peter J; Gorovits Elena L; Patti Joseph M Author Address: Inhibitex Inc, 9005 Westside Pkwy, Alpharetta, GA 30004 USA**USA Page 3

Author E-mail Address: jvernachio@inhibitex.com Journal: Antimicrobial Agents and Chemotherapy 50 (2): p 511-518 FEB 2006 2006

ISSN: 0066-4804

Document Type: Article Record Type: Abstract Language: English

Human immunoglobulin G recognizing fibrinogen-binding surface proteins is protective against both Staphylococcus aureus and Staphylococcus epidermidis infections in vivo

Abstract: A human donor-selected immunoglobulin G for intravenous injection (IGIV) product with elevated titers against the staphylococcal fibrinogen-binding MSCRAMM proteins ClfA and SdrG (INH-A21) was tested in vitro and in vivo. INH-A21 contained a significantly increased ability to inhibit the fibrinogen-binding activity of recombinant forms of both ClfA and SdrG. Evaluation of the opsonizing potential of INH-A21 was evaluated using fluorescently labeled bacteria; this... DESCRIPTORS:

Chemicals & Biochemicals: ...immunoglobulin G... ...SdrG;fibrinogen-binding surface proteins

>>>W: KWIC option is not available in file(s): 399 7/K/3 (Item 3 from file: 5) Links Fulltext available through: USPTO Full Text Retrieval Options Biosis Previews(R) (c) 2007 The Thomson Corporation. All rights reserved.

18192570 Biosis No.: 200500098483 A fibrinogen-binding protein of Staphylococcus lugdunensis

Author: Nilsson Martin; Bjerketorp Joakim; Guss Bengt; Frykberg Lars (Reprint) Author Address: Dept Microbiol, Swedish Univ Agr Sci, POB 7025, SE-75007, Uppsala, Sweden** Sweden

Author E-mail Address: lars.frykberg@mikrob.slu.se

Journal: FEMS Microbiology Letters 241 (1): p 87-93 December 1, 2004 2004

Medium: print ISSN: 0378-1097

Document Type: Article Record Type: Abstract Language: English

A fibrinogen-binding protein of Staphylococcus lugdunensis

Abstract: A gene called fbl, encoding a Staphylococcus lugdunensis fibrinogen -binding protein, was identified by phage display. The encoded protein, Fbl, is a member of the Sdr-family, a group of staphylococcal cell surface proteins containing a characteristic serine-aspartate repeat region. The fibrinogen-binding domain was mapped to 313 amino acids, and shows, 62% identity to the corresponding region in clumping factor (ClfA) from Staphylococcus aureus. Anti-serum against ClfA cross-reacted with Fbl, and blocked S. lugdunensis adherence to fibrinogen. Twelve clinical isolates of S. lugdunensis analysed by Southern blot all had an fbl-like...

DESCRIPTORS:

Chemicals & Biochemicals: fibrinogen;fibrinogen-binding protein

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7/K/4 (Item 4 from file: 5) Links

Fulltext available through: USPTO Full Text Retrieval Options

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Biosis No.: 200400395602

beta2-integrins mediate a novel form of chemoresistance in cycloheximide-induced U937 apoptosis

Author: Wu R-C; Wang Z (Reprint); Liu M-J; Chen D-F; Yue X-S

Author Address: Dept Biol Sci and Biotechnol, Tsing Hua Univ, Beijing, 100084,

China**China

Author E-mail Address: zwang@tsinghua.edu.cn Journal: CMLS Cellular and Molecular Life Sciences 61 (16): p 2071-2082 August

2004 2004 Medium: print ISSN: 1420-682X

Document Type: Article Record Type: Abstract Language: English

Abstract: ...leukaemic cell line U937, a novel form of chemoresistance, which we termed sudden drug resistance (SDR), was identified using Hoechst33258 staining, Western blott and DNA Ladder. CHXhigh (10-100 mug/ml)....inhibited by short-term preincubation with CHXlow (2.5 mug/ml). Unlike typical multidrug resistance, SDR is not caused by reduced drug accumulation or altered protein expression, and may be associated.....adhesion has been suggested to influence cell survival and prevent apoptosis. EDTA, or anti-CD18 monoclonal antibody, but not EGTA, acetylsalicylic acid or RGDS tetrapeptide, abrogated this homotypic aggregation and greatly increased CHX-induced apoptosis in a time-dependent manner, while fibrinogen and soluble intercellular adhesion molecule-1 exerted opposite effects. These results establish that beta2-integrin engagement is a key mediator of SDR, although it may be non-exclusive. This finding supplements the classical basis of chemoresistance and...

DESCRIPTORS:

Chemicals & Biochemicals: ...anti-CD18 monoclonal antibody.. Miscellaneous Terms: Concept Codes: sudden drug resistance {SDR}

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USPTO Full Text Retrieval Options

Biosis Previews(R)

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Biosis No.: 200300519934

Methicillin-resistant Staphylococcus aureus isolates carrying pls invade host cells less efficiently than pls-negative MRSA isolates.

Author: Sinha B (Reprint); Juuti K; Werbick C (Reprint); Kuusela P; Peters G (Reprint)

Author Address: Institute of Medical Microbiology, University Hospital of Muenster, Muenster, Germany**Germany

Journal: Abstracts of the General Meeting of the American Society for Microbiology 103 p B-207 2003 2003

Medium: cd-rom

Conference/Meeting: 103rd American Society for Microbiology General Meeting Washington, DC, USA May 18-22, 2003; 20030518
Sponsor: American Society for Microbiology
ISSN: 1060-2011 _(ISSN print)

Document Type: Meeting; Meeting Abstract

Record Type: Abstract

Language: English

Abstract: ...MSSA isolates. Pls (plasmin-sensitive) is a cell wall-anchored surface protein, belonging to the Sdr family of adhesins. Since adherence of pls-positive MRSA isolates to immobilized IgG, fibrinogen and Fn is reduced, we tested, whether this is also true for cellular invasiveness. Methods... **DESCRIPTORS:**

Chemicals & Biochemicals: IgG {immunoglobulin G...

KWIC option is not available in file(s): 399 >>>W: Page 5

sdrgantibody.txt 7/K/6 (Item 6 from file: 5) Links Biosis Previews(R) (c) 2007 The Thomson Corporation. All rights reserved. Biosis No.: 200300277832 17309113 MSCRAMM(R) protein mAb protects against S. epidermidis central venous catheter induced infection. Author: Vernachio J (Reprint); Bryant D (Reprint); Hall A (Reprint); Patel P (Reprint); Domanski P (Reprint); Syribeys P (Reprint); Gorovits E (Reprint) ; Wang J (Reprint); Robbins J (Reprint); Hutchins J (Reprint); Patti J (Reprint)
Author Address: Inhibitex, Inc., Alpharetta, GA, USA**USA
Journal: Abstracts of the Interscience Conference on Antimicrobial Agents and Chemotherapy 42 p 32 2002 2002 Medium: print Conference/Meeting: American Society for Microbiology (ASM) Annual Meeting on Infectious Disaease San Diego, CA, USA September 27-30, 2002; 20020927 Sponsor: American Society for Microbiology Document Type: Meeting; Meeting Abstract Record Type: Abstract Language: English Abstract: ...both a reduction in the incidence and severity of disease. We have demonstrated that a monoclonal antibody (mAb) against the MSCRAMM(R) protein, SdrG, inhibits the binding to human fibrinogen in vitro and also provides significant protection against methicillin resistant S. epidermidis (MRSE) challenge in... ...infection model. Methods: Clinical efficacy was evaluated in a rat model of CVC-associated infection. SdrG mAb 59-59 (n=10) and a control mAb (n=13) were administered IV. 24... ...were infected (13/13). Conclusions: These data clearly demonstrate that a single infusion with a SdrG mAb can significantly prevent catheter associated MRSE bacteremia and subsequent hematogenous dissemination to target organs.
DESCRIPTORS: Chemicals & Biochemicals: ...monoclonal antibodies... ...MSCRAMM protein monoclonal antibody KWIC option is not available in file(s): 399 7/K/7 (Item 7 from file: 5) Links Biosis Previews(R) (c) 2007 The Thomson Corporation. All rights reserved. 16972468 Biosis No.: 200200565979 Prevention of experimental Staphylococcus epidermidis (SE) endocarditis (IE) by passive immunotherapy with INH-A00021, a human IgG directed against staphylococcal fibrinogen-binding proteins Author: Kupferwasser L I (Reprint); Prater B; Wang J; Ruckstuhl M J; Lee K; Gast D; Adams D; Patti J M; Bayer A S (Reprint)
Author Address: Harbor-UCLA Res. and Ed. Inst., Torrance, CA, USA**USA
Journal: Abstracts of the Interscience Conference on Antimicrobial Agents and
Chemotherapy 41 p 278 2001 2001 Medium: print Conference/Meeting: 41st Annual Meeting of the Interscience Conference on

Abstract: Background: SE is a major cause of endovascular infections, utilizing adhesins such as the fibrinogen-binding adhesin, SdrG, to bind to sites of endovascular damage. Purpose: INH-A00021 is a donor-selected, plasma-derived Page 6

..SE) endocarditis (IE) by passive immunotherapy with INH-A00021, a human IgG

September 22-25,

Antimicrobial Agents and Chemotherapy Chicago, Illinois, USA

directed against staphylococcal fibrinogen-binding proteins

2001; 20010922

Record Type: Abstract Language: English

Document Type: Article; Meeting

sdrgantibody.txt hyperimmune globulin containing elevated levels of IgG against the staphylococcal fibrinogen-binding proteins, SdrG, and ClfA. This study evaluated the efficacy of INH-A00021 in attenuating experimental SE IE... ...p=0.0006). Also, the extent of bacteremia was significantly lower in animals receiving anti-SdrG, when compared to controls (p<0.01). Results of quantitative tissue cultures (mean log10CFU/g... DESCRIPTORS: Chemicals & Biochemicals: ...IgG {immunoglobulin G... ...SdrG--. ...fibrinogen-binding adhesion.....staphylococcal fibrinogen-binding proteins >>>W: KWIC option is not available in file(s): 399 7/K/8 (Item 8 from file: 5) Links Fulltext available through: USPTO Full Text Retrieval Options Biosis Previews(R) (c) 2007 The Thomson Corporation. All rights reserved. 15411492 Biosis No.: 200000129805 A bone sialoprotein-binding protein from Staphylococcus aureus: A member of the staphylococcal Sdr family Author: Tung Hui-shan; Guss Bengt; Hellman Ulf; Persson Lena; Rubin Kristofer; Ryden Cecilia (Reprint) Author Address: Department of Medical Biochemistry and Microbiology, Uppsala University, BMC, SE-751 23, Uppsala, Sweden**Sweden
Journal: Biochemical Journal 345 (3): p 611-619 Feb. 1, 2000 2000 Medium: print ISSN: 0264-6021 Document Type: Article Record Type: Abstract Language: English A bone sialoprotein-binding protein from Staphylococcus aureus: A member of the staphylococcal Sdr family Abstract: ...acids, called BSP-binding protein (Bbp), which displays similarity to recently described proteins of the Sdr family from S. aureus. SdrC, SdrD and SdrE encode putative cell-surface proteins with no described ligand specificity. Bbp also shows similarity to a fibrinogen -binding protein from S. epidermidis called Fbe. A serine-aspartic acid repeat sequence was found close to the cell-wall-anchoring Leu... protein bound radiolabelled native BSP, and inhibited the binding of radiolabelled BSP to staphylococcal cells. Serum from patients suffering from bone and joint infection contained antibodies that reacted with the fusion... DESCRIPTORS: Chemicals & Biochemicals: Sdr; >>>W: KWIC option is not available in file(s): 399 7/K/9 (Item 1 from file: 34) Links Fulltext available through: American Society for Microbiology custom link USPTO Full Text Retrieval Options SciSearch(R) Cited Ref Sci (c) 2007 The Thomson Corp. All rights reserved. 09577298 Genuine Article#: 423CT No. Refere No. References: 51 Expression of pls, a gene closely associated with the mecA gene of methicillin-resistant Staphylococcus aureus, prevents bacterial adhesion in vitro Author: Savolainen K (REPRINT); Paulin L; Westerlund-Wikstrom B; Foster TJ; Korhonen TK; Kuusela P Corporate Source: Univ Helsinki, Div Gen Microbiol, Dept Biosci, POB 56/FIN-00014
Helsinki//Finland/ (REPRINT); Univ Helsinki, Div Gen Microbiol, Dept Biosci, FIN-00014
Helsinki//Finland/; Univ Helsinki, Inst Biotechnol, FIN-00014 Helsinki//Finland/; Univ Helsinki, Haartman Inst, Dept Bacteriol & Immunol, FIN-00014 Helsinki//Finland/; Univ Helsinki, Cent Hosp, HUCH Lab Diagnost, Div Clin Microbiol, Helsinki//Finland/; Univ Dublin Trinity Coll, Moyne Inst Prevent Med, Dept Microbiol, Dublin 2//Ireland/ Journal: INFECTION AND IMMUNITY, 2001, V 69, N5 (MAY), P 3013-3020

ISSN: 0019-9567 Publication date: 20010500 Publisher: AMER SOC MICROBIOLOGY, 1752 N ST NW, WASHINGTON, DC 20036-2904 USA Language: English Document Type: ARTICLE (ABSTRACT AVAILABLE) Abstract: ...distinct repeat regions, one of which was a serine-aspartate repeat characteristic of the Clf-Sdr family of surface proteins in staphylococci, The lengths of the repeat regions varied in different....digested DNA. A pls mutant constructed by allele replacement adhered well to immobilized fibronectin and immunoglobulin e, in contrast to the parental strain, suggesting that Pls could have

identifiers-- ...FIBRINOGEN-BINDING PROTEIN; NUCLEOTIDE-SEQUENCE; CLUMPING FACTOR; INSERTIONAL INACTIVATION; ESCHERICHIA-COLI; REPEAT REGION; FIBRONECTIN; CLONING; DNA

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SciSearch(R) Cited Ref Sci
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02147492 Genuine Article#: KE378 No. References: 43
INHIBITION OF PLATELET-ADHESION TO FIBRIN(OGEN) IN FLOWING WHOLE-BLOOD BY

ARG-GLY-ASP AND FIBRINOGEN GAMMA-CHAIN CARBOXY TERMINAL PEPTIDES

Author: HANTGAN RR; ENDENBURG SC; CAVERO I; MARGUERIE G; UZAN A; SIXMA JJ; DEGROOT

Corporate Source: WAKE FOREST UNIV,BOWMAN GRAY SCH MED,DEPT BIOCHEM,MED CTR BLVD/WINSTON SALEM//NC/27157; UNIV UTRECHT,DEPT HEMATOL/UTRECHT/NETHERLANDS/; RHONE POULENC RORER RD/VITRY//FRANCE/; INSERM,U127,HEMATOL LAB/GRENOBLE//FRANCE/Journal: THROMBOSIS AND HAEMOSTASIS , 1992 , V 68 , N6 (DEC 7) , P 694-700 ISSN: 0340-6245

Language: ENGLISH Document Type: ARTICLE (Abstract Available)...OF PLATELET-ADHESION TO FIBRIN(OGEN) IN FLOWING WHOLE-BLOOD BY ARG-GLY-ASP AND FIBRINOGEN GAMMA-CHAIN CARBOXY TERMINAL PEPTIDES Abstract: We have employed synthetic peptides with sequences corresponding to the

Abstract: We have employed synthetic peptides with sequences corresponding to the integrin receptor-recognition regions of fibrinogen as inhibitors of platelet aggregation and adhesion to fibrinogen and fibrin-coated surfaces in flowing whole blood, using a rectangular perfusion chamber at wall.....1,300 s-1. D-RGDW caused substantial inhibition of platelet aggregation and adhesion to fibrinogen and fibrin at both shear rates, although it was least effective at blocking platelet adhesion....300 s-1. RGDS was a weaker inhibitor, and produced a biphasic dose-response curve; SDRG was inactive. HHLGGAKQAGDV partially inhibited platelet aggregation and adhesion to fibrin(ogen) at both shear ...

Identifiers-- ...GLYCOPROTEIN-IIB-IIIA; VONWILLEBRAND-FACTOR; MONOCLONAL -ANTIBODIES; ARTIFICIAL SURFACES; BINDING; RECEPTOR; SUBENDOTHELIUM; COMPLEX; CELLS;

FIBRONECTIN
Research Fronts: 91-2339 004 (PLATELET GLYCOPROTEIN-IIB-IIIA COMPLEX; FIBRINOGEN
RECEPTOR ANTAGONIST; ANTIPLATELET ARG-GLY-ASP-CONTAINING PEPTIDE; SNAKE-VENOM
PROTEIN ECHISTATIN)

91-5876 001...

Nilsson, M.
Department of Microbiology, Swedish University of Agricultural Sciences, Box 7025, S-750 07 Uppsala, Sweden.
Acta Universitatis Agriculturae Sueciae - Agraria (265): p.115
Publication Year: 2001

ISSN: 1401-6249 Publisher: Sveriges Lantbruksuniversitet (Swedish University of Agricultural Uppsala , Sweden Sciences) ISBN: 91-576-5791-2 Language: English Record Type: Abstract Document Type: Thesis Fibrinogen - and von Willebrand factor-binding proteins in staphylococci. ... genes, isolated from coagulase-negative staphylococci (CoNS) associated with human infections, and their corresponding proteins. Fbe and Fbl are the main fibrinogen (Fg)-binding proteins of Staphylococcus epidermidis and S. lugdunensis, respectively. Both proteins are members of the Sdr (SD-repeat containing protein) family, a subgroup of cell surface proteins in staphylococci with a... ... less perfect, tandemly repeated serine and aspartate residues. Sequence comparisons in the binding regions between Fbe and Fbl revealed low mutual similarity. However, Fbl is relatively conserved (63% identity) in the binding region compared to clumping factor A (ClfA), the prototype Sdr protein from S. aureus. The third gene, vWbl, encodes a putative von Willebrand factor (vWf...... an overall organization, that is characteristic for cell surface proteins in staphylococci. The importance of Fbe, Fbl and vWbl for their respective organisms during the infection process is not known, but ... to extracellular matrix or plasma-coated biomaterials. Separate recombinant constructs, comprising the binding regions of Fbe and Fbl or separate antibodies directed against the binding regions of the proteins, were able......
the adherence of S. epidermidis and S. lugdunensis, respectively, to immobilized
Fg. The presence of fbe, fbl and vwbl genes is very common in clinical isolates of
the respective species. In..... these experiments, vwbp immobilized on a
Sepharose-column was used to purify vwf from human serum. The gene vwb was present in all tested strains of S. aureus. Descriptors: ...fibrinogen; >>>W: KWIC option is not available in file(s): 399 7/K/12 (Item 1 from file: 71) Links Fulltext available through: USPTO Full Text Retrieval Options ELSEVIER BIOBASE (c) 2007 Elsevier B.V. All rights reserved. 02747070 2004224218 betaSUB2-integrins mediate a novel form of chemoresistance in cycloheximide-induced U937 apoptosis Wu R.-C.; Wang Z.; Liu M.-J.; Chen D.-F.; Yue X.-S. Address: Z. Wang, Dept. of Biol. Sci. and Biotech., Tsinghua University, Beijing, China Email: zwang@tsinghua.edu.cn Journal: Cellular and Molecular Life Sciences, 61/16 (2071-2082), 2004, Switzerland CODEN: CMLSF ISSN: 1420-682X Document Type: Article Languages: English Summary Languages: English No. of References: 48 **DESCRIPTORS:** Apoptosis; Cycloheximide; U937 cell; betaSUB2-integrin; Drug resistance; PI-3K CLASSIFICATION CODE AND DESCRIPTION: Modlecular Sequence Databank Number: 87.2.1.5 - CANCER RESEARCH / TUMOUR BIOLOGY / Cellular Biology and Biochemistry / Immortalisation, senescence and apoptosis 87.4.1.15 - CANCER RESEARCH / TREATMENT / Chemotherapy / Resistance 87.4.11 - CANCER RESEARCH / TREATMENT / Treatment Monitoring and Evaluation ...leukaemic cell line U937, a novel form of chemoresistance, which we termed sudden drug resistance (SDR), was identified using Hoechst33258 staining, Western blott and

sdrgantibody.txt DNA Ladder. CHXSUPhigh (10-100 mug/ml... ...inhibited by short-term preincubation with CHXSUBlow (2.5 mug/ml). Unlike typical multidrug resistance, SDR is not caused by reduced drug accumulation or altered protein expression, and may be associated...
...has been suggested to influence cell survival and prevent apoptosis. EDTA, or
anti-CD 18 monoclonal antibody, but not EGTA, acetylsalicylic acid or RGDS
tetrapeptide, abrogated this homotypic aggregation and greatly increased CHX-induced
apoptosis in a time-dependent manner, while fibrinogen and soluble intercellular adhesion molecule-1 exerted opposite effects. These results establish that betaSUB2-integrin engagement is a key mediator of SDR, although it may be non-exclusive. This finding supplements the classical basis of chemoresistance and... >>W: KWIC option is not available in file(s): 399 7/K/13 (Item 1 from file: 155) Links >>>W: Fulltext available through: USPTO Full Text Retrieval Options MEDLINE(R) (c) format only 2007 Dialog. All rights reserved. 15223135 PMID: 15583173 Protein FOG--a streptococcal inhibitor of neutrophil function. Johansson Helena M; Morgelin Matthias; Frick Inga-Maria
Department of Cell and Molecular Biology, Section for Clinical and Experimental
Infectious Medicine, BMC, B14, Lund University, S-221 84 Lund, Sweden.
Microbiology (Reading, England) (England) Dec 2004, 150 (Pt 12) p4211-21,
ISSN: 1350-0872--Print Journal Code: 9430468 Publishing Model Print Document type: Journal Article; Research Support, Non-U.S. Gov't Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed ...of group G streptococci (GGS) form aggregates when grown in vitro. Aggregating strains interact with fibrinogen, and this study reports the isolation of a novel self-associating and fibrinogen-binding protein of GGS, denoted process. Sequencing of the fog gene revealed structural similarity....of GGS express protein G, a protein known to interact with the constant region of immunoglobulin G and albumin. Surprisingly, a clinical isolate expressing protein G, but lacking protein FOG, was... ...negative strain from being killed. The antibactericidal property of protein FOG is dependent on its fibrinogen-binding activity. Thus, in plasma, FOG precipitates fibrinogen, and when added to whole blood, protein FOG triggers the formation of visible aggregates comprising fibrinogen and neutrophils that are disabled in their killing of the bacteria. Moreover, the results emphasize.. Descriptors: *Bacterial Proteins--metabolism--ME; *Blood--microbiology--MI; *Carrier Proteins--metabolism--ME; *Fibrinogen--metabolism--ME; *Neutrophils --immunology--IM; *Streptococcus--growth and development--GD Chemical Name: Bacterial Proteins, Carrier Proteins, Fbe protein, bacteria; Fibrinogen >>>W: KWIC option is not available in file(s): 399 7/K/14 (Item 1 from file: 393) Links Beilstein Database - Abstracts (c) 2007 Beilstein GmbH. All rights reserved. Beilstein Abstract Id: 6577659 Title: Human Immunoglobulin G Recognizing Fibrinogen-Binding Surface Proteins Is Protective against both Staphylococcus aureus and Staphylococcus epidermidis Infections In Vivo Document Type: Journal Record Type: Abstract Author: Vernachio, John H.; Bayer, Arnold S.; Ames, Brenda; Bryant, Dawn; Prater, Bradley D.; Syribeys, Peter J.; Gorovits, Elena L.; Patti, Joseph M. Citation: Antimicrob. Agents & Chemother. (2006) Series: 50-2, 511 - 518 CODEN:

Page 10

AMACCQ Language: English

Abstract Language: English

Title: Human Immunoglobulin G Recognizing Fibrinogen-Binding Surface Proteins Is Protective against both Staphylococcus aureus and Staphylococcus epidermidis Infections In Vivo Abstract: A human donor-selected immunoglobulin G for intravenous injection (IGIV) product with elevated titers against the staphylococcal fibrinogen-binding MSCRAMM proteins ClfA and SdrG (INH-A2Ĭ) was tested in vitro and in vivo. INH-A21 contained a significantly increased ability to inhibit the fibrinogen-binding activity of recombinant forms of both ClfA and SdrG. Evaluation of the opsonizing potential of INH-A21 was evaluated using fluorescently labeled bacteria; this... >>W: KWIC option is not available in file(s): 399 7/K/15 (Item 2 from file: 393) Links Beilstein Database - Abstracts (c) 2007 Beilstein GmbH. All rights reserved. Beilstein Abstract Id: 6471865 Title: beta 2 -integrins mediate a novel form of chemoresistance in cycloheximide-induced U937 apoptosis Document Type: Journal Record Type: Abstract Author: Wu, R.-C.; Wang, Z.; Liu, M.-J.; Chen, D.-F.; Yue, X.-S. Citation: Cell. Mol. Life Sci (2004) Series: 61-16, 2071 - 2082 CODEN: CMLSFI Language: English Abstract Language: English Keywords: apoptosis; cycloheximide; U937 cell; Beta 2 -integrin; drug resistance; PI-3K Abstract: ... leukaemic cell line U937, a novel form of chemoresistance, which we termed sudden drug resistance (SDR), was identified using Hoechst33258 staining, Western blott and DNA Ladder. CHX high (10-100 mu... short-term preincubation with CHX low (2.5 mu g/ml). Unlike typical multidrug resistance, SDR is not caused by reduced drug accumulation or altered protein expression, and may be associated... adhesion has been suggested to influence cell survival and prevent apoptosis. EDTA, or anti-CD18 monoclonal antibody, but not EGTA, acetylsalicylic acid or RGDS tetrapeptide, abrogated this homotypic aggregation and greatly increased CHX-induced apoptosis in a time-dependent manner, while fibrinogen and soluble intercellular adhesion molecule-1 exerted opposite effects. These results establish that beta 2 -integrin engagement is a key mediator of SDR, although it may be non-exclusive. This finding supplements the classical basis of chemoresistance and... KWIC option is not available in file(s): 399 7/K/16 (Item 1 from file: 399) Links CA SEARCH(R) (c) 2007 American Chemical Society. All rights reserved. CA: 144(19)348882p 144348882 **PATENT** Immunogenic composition comprising a mixture of staphylococcal antigens and uses as vaccines Inventor (Author): Castado, Cindy; Lecrenier, Nicolas Pierre Fernand; Neyt, Cecile Anne; Poolman, Jan Location: Belg. Assignee: GlaxoSmithKline Biologicals S.A. Patent: PCT International; WO 200632472 A2 Date: 20060330 Application: WO 2005EP10184 (20050920) *GB 200421082 (20040922) *GB 200421078 (20040922) *GB 200421081 (20040922) *GB 200421079 (20040922) *GB 20053143 (20050215) Pages: 132 pp. CODEN: PIXXD2 Language: English Patent Classifications: A61K-000/A Class: Designated Countries: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY; BZ; CA; CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD; GE; GH; GM; HR; Page 11

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sdrgantibody.txt
HU; ID; IL; IN; IS; JP; KE; KG; KM; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; LY; MA;
MD; MG; MK; MN; MW; MX; MZ; NA; NG; NI; NO; NZ; OM; PG; PH; PL; PT; RO; RU; SC; SD;
SE; SG; SK; SL; SM; SY; TJ; TM; TN; TR; TT; TZ; UA; UG; US; UZ; VC; VN; YU; ZA; ZM Designated Regional: AT; BE; BG; CH; CY; CZ; DE; DK; EE; ES; FI; FR; GB; GR; HU; IE; IS; IT; LT; LU; LV; MC; NL; PL; PT; RO; SE; SI; SK; TR; BF; BJ; CF; CG; CI; CM; GA; GN; GQ; GW; ML; MR; NE; SN; TD; TG; BW; GH; GM; KE; LS; MW; MZ; NA; SD; SL; SZ; TZ; UG; ZM; ZW; AM; AZ; BY; KG; KZ; MD; RU; TJ; TM
>>>W: KWIC option is not available in file(s): 399
 7/K/17 (Item 2 from file: 399) Links
CA SEARCH(R)
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                       CA: 141(21)348821f
                                                            PATENT
Staphylococcus epidermidis-derived hyperimmune serum reactive antigens and encoding
nucleic acids for diagnosis and treatment of bacterial infection and for antagonist
Inventor (Author): Meinke, Andreas; Min, Bui Duc; Nagy, Eszter
Location: Austria
Assignee: Intercell AG
Patent: PCT International; WO 200487746 A2 Date: 20041014 Application: WO 2004EP3398 (20040331) *EP 2003450078 (20030331)
                                                                   Date: 20041014
Pages: 196 pp.
CODEN: PIXXD2
Language: English
Patent Classifications:
   Class:
               C07K-014/00A
Designated Countries: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BW; BY; BZ; CA;
CH; CN; CO; CR; CU; CZ; DE; DK; DM; DZ; EC; EE; EG; ES; FI; GB; GD; GE; GH; GM; HR; HU; ID; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; MA; MD; MG; MK; MN; MX; MZ; NA; NI; NO; NZ; OM; PG; PH; PL; PT; RO; RU; SC; SD; SE; SG; SK; SL; SY; TJ; TM; TN; TT; TZ; UA; UG; US; UZ; VC; VN; YU; ZA; ZM; ZW
Designated Regional: BW; GH; GM; KE; LS; MW; MZ; SD; SL; SZ; TZ; UG; ZM; ZW; AM; AZ;
BY; KG; KZ; MĎ; RU; TJ; TM; AT; BE; BG; CH; CY; CZ; DE; DK; EE; ES; FI; FR; GB; GR;
HU; IE; IT; LU; MC; NL; PL; PT; RO; SE; SI; SK; TR; BF; BJ; CF; CG; CI; CM; GA; GN;
GQ; GW; ML; MR; NE; SN; TD; TG
         KWIC option is not available in file(s): 399
 7/K/18 (Item 3 from file: 399) Links
CA SEARCH(R)
(c) 2007 American Chemical Society. All rights reserved.
139259972
                       CA: 139(17)259972x
                                                            PATENT
Monoclonal and polyclonal antibodies recognizing coagulase-negative staphylococcal
Inventor (Author): Patti, Joseph M.; Hutchins, Jeff T.; Hall, Andrea; Domanski,
Paul; Patel, Pratisksha; Hook, Magnus; Robbins, Jeff; Vernachio, John; Bowden, Maria
G.
Location: USA
Assignee: Inhibitex, Inc.; The Texas A & M University System Patent: PCT International; WO 200376470 A1 Date: 200309 Application: WO 2003US6415 (20030305) *US PV361324 (20020305)
                                                                   Date: 20030918
Pages: 72 pp.
CODEN: PIXXD2
Language: English
Patent Classifications:
               CO7K-016/00A; CO7K-001/00B; CO7K-002/00B; CO7H-021/04B; A61K-039/395B;
A61K-039/40B; A61K-039/00B; A61K-039/09B; A61K-039/085B
Designated Countries: AE; AG; AL; AM; AT; AU; AZ; BA; BB; BG; BR; BY; BZ; CA; CH;
CN; ČO; CR; CU; CZ; DE; ĎK; ĎM; ĎZ; ÉC; ÉE; ÉS; FI; ŚB; ŚD; ŚE; ŚH; ŚM; ĤR; ĤU; ÍD; IL; IN; IS; JP; KE; KG; KP; KR; KZ; LC; LK; LR; LS; LT; LU; LV; MA; MD; MG; MK; MN;
                                                     Page 12
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sdrgantibody.txt
MW; MX; MZ; NO; NZ; OM; PH; PL; PT; RŎ; RU; SĆ; SD; SE; SG; SK; SL; TJ; TM; TN; TR;
TT; TZ; UA; UG; UZ; VC; VN; YU; ZA; ZM; ZW; AM; AZ; BY; KG; KZ; MD; RU; TJ; TM
Designated Regional: GH; GM; KE; LS; MW; MZ; SD; SL; SZ; TZ; UG; ZM; ZW; AT; BE; BG; CH; CY; CZ; DE; DK; EE; ES; FI; FR; GB; GR; HU; IE; IT; LU; MC; NL; PT; RO; SE; SI; SK; TR; BF; BJ; CF; CG; CI; CM; GA; GN; GQ; GW; ML; MR; NE; SN; TD; TG
>>>W:
        KWIC option is not available in file(s): 399
 7/K/19 (Item 1 from file: 357)
                                        Links
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    Fulltext available through:
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0420861 DBA Accession No.: 2007-06799
A panel of monoclonal antibodies recognizing the Staphylococcus epidermidis fibrinogen-binding MSCRAMM SdrG hybridoma cell culture for recombinant monoclonal
antibody production
Author: HALL AE; PATEL PR; DOMANSKI PJ; PRATER BD; GOROVITS EL; SYRIBEYS PJ;
VERNACHIO JH; PATTI JM; HUTCHINS JT
Corporate Affiliate: Inhibitex Inc
Corporate Source: Hutchins JT, Inhibitex Inc, 9005 Westside Pkwy, Alpharetta, GA
30004 USA
Journal: HYBRIDOMA ( 26, 1, 28-34 )
                                                 2007
ISSN: 1554-0014
Language: English
A panel of monoclonal antibodies recognizing the Staphylococcus epidermidis
fibrinogen-binding MSCRAMM SdrG hybridoma cell culture for recombinant monoclonal
antibody production
Abstract: ...stage contributing to the pathogenesis of this bacteria is the initial
adherence to host tissue. SdrG is a cell-wall-anchored fibrinogen-binding adhesin of S. epidermidis that has been shown to be necessary for bacterial binding to fibrinogen-coated foreign bodies, such as catheters. Here we report the generation and characterization of a panel of monoclonal antibodies (MAbs) directed against this S. epidermidis virulence factor. Through the use of multiple in.....that may
prove to be beneficial in studies that address the precise biologic role of SdrG. (7
Descriptors: Staphylococcus epidermidis fibrinogen-binding MSCRAMM SdrG -specific
recombinant monoclonal antibody prep., purification, characterization,
plasmid-mediated gene transfer, expression in Lactococcus lactis, hybridoma, mouse
immunization...
        KWIC option is not available in file(s): 399
 7/K/20 (Item 2 from file: 357)
                                        Links
                                         USPTO Full Text Retrieval Options
    Fulltext available through:
Derwent Biotech Res.
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0358809 DBA Accession No.: 2005-04513
A fibrinogen-binding protein of Staphylococcus lugdunensis identification and production of a recombinant fibrinogen binding protein from Staphylococcus
lugdunensis using phage display and recombinant technology
Author: NILSSON M; BJERKETORP J; GUSS B; FRYKBERG L
Corporate Affiliate: Swedish Univ Agr Sci
Corporate Source: Frykberg L, Swedish Univ Agr Sci, Dept Microbiol, POB 7025,
SE-75007 Uppsala, Sweden
Journal: FEMS MICROBIOLOGY LETTERS (241, 1, 87-93)
                                                                      2004
ISSN: 0378-1097
Language: English
A fibrinogen-binding protein of Staphylococcus lugdunensis identification and
production of a recombinant fibrinogen binding protein from Staphylococcus
lugdunensis using phage display and recombinant technology
Abstract: AUTHOR ABSTRACT - A gene called fbl, encoding a Staphylococcus lugdunensis
                                                Page 13
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fibrinogen-binding protein, was identified by phage display. The encoded protein, Fbl, is a member of the Sdr-family, a group of staphylococcal cell surface proteins containing a characteristic serine-aspartate repeat region. The fibrinogen-binding domain was mapped to 313 amino acids, and shows, 62% identity to the corresponding region in clumping factor (ClfA) from Staphylococcus aureus. Anti- serum against ClfA cross-reacted with Fbl, and blocked S. lugdunensis adherence to fibrinogen. Twelve clinical isolates of S. lugdunensis analysed by Southern blot all had an fbl-like...

Descriptors: Staphylococcus lugdunensis, recombinant fibrinogen binding protein, prep, isol., characterization, phage display, fbl gene identification, Southern blot bacterium surface display...

>>>: KWIC option is not available in file(s): 399. 7/K/21 (Item 3 from file: 357) Links Derwent Biotech Res. (c) 2007 The Thomson Corp. All rights reserved. 0324377 DBA Accession No.: 2003-25518 PATENT New antibody recognizing a Staphylococcus epidermidis protein comprising SdrG N1N2N3, SdrG N2N3 or SdrGTR2 useful for preparing a composition for treating or preventing a coagulase-negative Staphylococcal infection chimeric antibody, humanized antibody, monoclonal antibody and single chain antibody production for vaccine, gene therapy and therapy Author: PATTI J M; HUTCHINS J T; HALL A; DOMANSKI P; PATEL P; HOOK M; ROBBINS J; VERNACHIO J; BOWDEN M G Patent Assignee: INHIBITEX INC; UNIV TEXAS A and M SYSTEM 2003 Patent Number: WO 200376470 Patent Date: 20030918 WPI Accession No.: 2003-722324 (200368)Priority Application Number: US 361324 Application Date: 20020305 National Application Number: WO 2003US6415 Application Date: 20030305 Language: English New antibody recognizing a Staphylococcus epidermidis protein comprising SdrG N1N2N3, SdrG N2N3 or SdrGTR2 useful for preparing a composition for treating or preventing a coagulase-negative Staphylococcal infection chimeric antibody, humanized antibody, monoclonal antibody and single chain antibody production for vaccine, gene therapy and therapy Abstract: DERWENT ABSTRACT: NOVELTY - An isolated antibody (I) that recognizes a protein from Staphylococcus epidermidis comprising SdrG N1N2N3, SdrG N2N3 or SdrGTR2, is new. DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for: (1) isolated....treating or preventing a coagulase-negative Staphylococcal infection; (7) an isolated S. epidermidis protein comprising SdrG N1N2N3, SdrG N2N3 or SdrGTR2; (8) eliciting an immunogenic reaction in a human or animal; (9) a... or SdrGTR2; (8) eliciting an immunogenic reaction in a human or animal; (9) a... ...epidermidis protein; and (10) an isolated nucleic acid sequence encoding an S. epidermidis protein comprising SdrG N1N2N3, SdrG N2N3 or SdrGTR2. BIOTECHNOLOGY - Preferred Antibody: The antibody (I) is selected from chimeric, murine, humanized or human monoclonal antibodies. (I) is a single chain monoclonal antibody. (I) binds to the S. epidermidis SdrG protein. (I) recognizes an amino acid sequence selected from the fully defined sequence comprising 560.......or 951 (S6) base pairs, respectively, as given in the specification. (II) and (III) are monoclonal antibodies. Preferred Kit: The kit comprises means for detecting binding by the antibody, which comprises a detectable label linked to the antibody. Preferred Protein: The isolated S. epidermidis protein comprising SdrG N1N2N3. SdrG N2N3 or Protein: The isolated S. epidermidis protein comprising SdrG N1N2N3, SdrG N2N3 or SdrGTR2 comprises an amino acid sequence selected from S1-S3 encoded by a ...Staphylococcal infection in a human or an animal and inhibits binding of Staphylococcal bacteria to fibrinogen, useful for preparing a composition for treating or preventing a coagulase-negative Staphylococcal infection. The monoclonal antibodies (II) and (III) comprising 1092 amino acids and 549 amino acids, respectively are also.....for treating or preventing a coagulase-negative Staphylococcal infection. An isolated S. epidermidis protein comprising SdrG N1N2N3, an Edge CR2 is administrated to a human on a primal to aligning the same and SdrG N2N3 or SdrGTR2 is administered to a human or animal to elicit an immune reaction.. Descriptors: Staphylococcus sp. epidermis-specific chimeric antibody, humanized

antibody, monoclonal antibody, single chain antibody prep., appl. vaccine, gene therapy, therapy antibody engineering antibody engineering protein... >>>w: KWIC option is not available in file(s): 399 7/K/22 (Item 1 from file: 149) Links TGG Health&wellness DB(SM) (c) 2007 The Gale Group. All rights reserved. Supplier Number: 73924880 (USE FORMAT 7 OR 9 FOR FULL TEXT) Whole genome sequencing of meticillin-resistant Staphylococcus aureus. Kuroda, Makoto; Ohta, Toshiko; Uchiyama, Ikuo; Baba, Tadashi; Yuzawa, Harumi; Kobayashi, Ichizo; Cui, Longzhu; Oguchi, Akio; Aoki, Ken-ichi; Nagai, Yoshimi; Lian, JianQi; Ito, Teruyo; Kanamori, Mutsumi; Matsumaru, Hiroyuki; Maruyama, Atsushi; Murakami, Hiroyuki; Hosoyama, Akira; Mizutani-Ui, Yoko; Takahashi, Noriko K; Sawano, Toshihiko; Inoue, Ryu-ichi; Kaito, Chikara; Sekimizu, Kazuhisa; Hirakawa, Hideki; Kuhara, Satoru; Goto, Susumu; Yabuzaki, Junko; Kanehisa, Minoru; Yamashita, Atsushi; Oshima, Kenshiro; Furuya, Keiko; Yoshino, Chie; Shiba, Tadayoshi; Hattori, Masahira; Ogasawara, Naotake; Hayashi, Hideo; Hiramatsu, Keiichi
The Lancet, 357, 9264, 1225 April 21, 2001 Publication Format: Magazine/Journal; Refereed ISSN: 0099-5355 Language: English Record Type: Fulltext; Abstract Target Audience: Professional Word Count: 10338 Line Count: 01076 Descriptors: Staphylococcus aureus--Genetic aspects; Genetic recombination--Physiological aspects; Drug resistance--Genetic aspects File Segment: HI File 149 ...wall sorting signal in N315 and Mu50 genomes (table 4). These include seven known adhesins: fibrinogen-binding proteins ClfA, ClfB, and SdrC-D-E, and fibronectin-binding proteins FnbA and FnbB... ...to form clusters at several loci in the genome rather than being randomly distributed. The fibrinogen-binding proteins are characteristic in their possession of serine-aspartate repeats that precede the LPXTG... ...similar to Streptococcus pyogenes myosin-reactive protein,(30) which is known to react with the serum of patients with acute rheumatic fever. The other three open reading frames (SA1751, SA0841, and... Ryden C. A bone sialoprotein-binding protein from Staphylococcus aureus: a member of the staphylococcal Sdr family. Biochem J 2000; 345: 611-19.

(28) Schneewind O, Model P, Fischetti VA. Sorting...
haemolysin

SAS065 Probable haemolysin SA1973 SA2207, 2208, 2209 g-haemolysin components Ser-Asp rich fibrinogen-binding proteins SA0742, 2423 0519, 0520, 0521 SA0587 Probable adhesin Possible extracellular matrix SA0744, 0745 binding proteins
Possible fibrinogen_binding proteins SA1000, 1003, 1004 SA1267, 1268 Probable extracellular matrix binding proteins Elastin-binding protein...

sdrgantibody.txt

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sdrgantibody.txt
...adhesion proteins
                                SA2459, 2460, 2461, 2462
    Others
    Myosin-crossreactive MHC class
                                                 SA0102
     II-like protein
     Immunoglobulin G binding protein A
                                                 SA0107
    Possible siderophore biosynthesis
                                                 SA0116, 0117
     proteins
    Probable capsular polysaccharide
                                                 SA0126...
```

d-haemolysin h1d

Probable haemolysin

g-haemolysin components hlgA, hlgC, hlgB

Ser-Asp rich fibrinogen-binding proteins clfA, clfB

sdrC, sdrD, sdrE

Probable adhesin

Possible extracellular matrix

binding proteins

Possible fibrinogen-binding proteins

Probable extracellular matrix ebhA, ebhB

ebpS

binding proteins
Elastin-binding protein
Fibronectin-binding...

...Intercellular adhesion proteins icaA, icaD, icaB, icaC

Others Myosin-crossreactive MHC class ÍI-like protein Immunoglobulin G binding protein A

spa Possible siderophore biosynthesis

proteins Probable capsular polysaccharide

synthesis proteins

Capsular...

SaPIn1/SaPIml ...toxin 1 d-haemolysin

Probable haemolysin g-haemolysin components

Adhesins

Ser-Asp rich fibrinogen-binding proteins

Probable adhesin

Possible extracellular matrix

binding proteins
Possible fibrinogen-binding proteins
Probable extracellular matrix

binding proteins Elastin-binding protein

Fibronectin-binding proteins

Intercellular adhesion proteins

Others

Myosin-crossreactive MHC class

II-like protein

Immunoglobulin G binding protein A Possible siderophore biosynthesis

proteins

Probable capsular polysaccharide

synthesis proteins

Capsular polysaccharide...

...cells

Probable haemolysin Unknown

g-haemolysin components Destruction of blood cells

Adhesins

Ser-Asp rich fibrinogen

-binding proteins Cellular adhesion onto

host tissues

Probable adhesin Cellular adhesion onto host tissues

Possible extracellular matrix Cellular adhesion onto

binding proteins
Possible fibrinogen-binding proteins
Probable extracellular matrix host tissues

Unknown Unknown

binding proteins Elastin-binding protein Cellular adhesion onto...

...infected tissues

Others

Myosin-crossreactive MHC class Potential immune disorder

in host

II-like protein Immunoglobulin

G binding protein A Potential immune disorder

in host

Possible siderophore biosynthesis Iron uptake

proteins...